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## (57) Abstract

This invention provides methods for treating a mycobacterial infection by administering to an animal a pharmaceutical composition containing a compound having the formula R-SO<sub>n</sub>-Z-CO-Y, where R is an alkyl group having 6-20 carbons; Z is a radical selected from -CH<sub>2</sub>-, -O-, and -NH-, two of these radicals coupled together, or -CH<sub>2</sub>-CH<sub>2</sub>-; Y is -NH<sub>2</sub>, O-CH<sub>2</sub>-C<sub>6</sub>H<sub>5</sub>, -CO-CO-O-CH<sub>3</sub>, or O-CH<sub>3</sub>; and n is 1 or 2. It has been discovered that these compounds inhibit growth of microbial cells which synthesize  $\alpha$ -substitued,  $\beta$ -hydroxy fatty acids, particularly conynemycolic acid, nocardic acid, and mycolic acid. These compounds may be used to inhibit growth of mycobacterial cells, such as Mycobacterium tuberculosis, drug-resistant M. tuberculosis, M. avium intracellulare, and M. leprae.